## **AMENDMENTS TO THE CLAIMS**

This listing of claims will replace all prior versions, and listings, of claims in the application.

- 1. (Original) A method of diagnosing a patient as having a neoplasia, said method comprising detecting an endocan nucleic acid molecule or polypeptide in a patient sample, wherein detection of an endocan nucleic acid or polypeptide indicates that said patient has a neoplasia.
- 2. (Original) The method of claim 1, wherein said patient sample is a blood sample.
- 3. (Original) The method of claim 1, wherein said patient sample is a tissue sample.
- 4. (Original) The method of claim 1, wherein said method comprises detecting an increase in the level of expression of an endocan polypeptide in a patient sample relative to the level of endocan polypeptide present in a corresponding control sample from a normal individual.
- 5. (Original) The method of claim 4, wherein said level of expression is determined in an immunological assay.
- 6. (Original) The method of claim 5, wherein said level of expression is detected by ELISA.

- 7. (Original) The method of claim 1, wherein said patient is asymptomatic.
- 8. (Original) A method of assessing the responsiveness of a neoplasm to a treatment regimen, said method comprising determining the level of an endocan nucleic acid or polypeptide in a patient sample relative to the level in a reference sample, wherein an alteration in said nucleic acid or polypeptide level in said patient sample indicates the responsiveness of said neoplasm to a treatment regimen.
- 9. (Original) The method of claim 8, wherein said reference sample is derived from a healthy individual.
- 10. (Original) The method of claim 8, wherein said patient is being treated for a neoplasm.
- 11. (Original) The method of claim 8, wherein said reference sample is obtained from said patient prior to or during the course of said treatment regimen.
- 12. (Original) The method of claim 8, wherein said patient sample is a blood sample.
- 13. (Original) The method of claim 8, wherein said patient sample is a tissue sample.
- 14. (Original) The method of claim 8, wherein said alteration is an increase, and said increase indicates a decreased responsiveness of said neoplasm to a treatment regimen.

- 15. (Original) The method of claim 8, wherein said alteration is a decrease, and said decrease indicates an increased responsiveness of said neoplasm to a treatment regimen.
- 16. (Original) The method of claim 8, wherein said neoplasm is a renal cell carcinoma, a lung cancer, a glioma, or a breast carcinoma.
- 17. (Original) A method of determining the prognosis of a patient having a neoplasm, said method comprising detecting an alteration in the level of an endocan nucleic acid molecule or polypeptide in a patient sample relative to the level in a reference sample, wherein an alteration indicates the prognosis of said patient.
- 18. (Original) The method of claim 17, wherein said reference sample is obtained from said patient prior to or during the course of said treatment regimen.
- 19. (Original) The method of claim 17, wherein said alteration is an increase, and said increase indicates a poor prognosis.
- 20. (Original) The method of claim 17, wherein said alteration is a decrease, and said decrease indicates a good prognosis.
- 21. (Original) The method of claim 17, wherein said patient sample is a tissue sample or a blood sample.
- 22. (Original) The method of claim 17, wherein said level of expression is determined in an immunological or enzymatic assay.

- 23. (Original) The method of claim 17, wherein said neoplasm is a renal cell carcinoma, a lung cancer, a glioma, or a breast carcinoma.
- 24. (Original) A diagnostic kit for the detection of a neoplasm in a patient comprising an endocan nucleic acid or amino acid sequence, or a fragment thereof.
- 25. (Original) A diagnostic kit for the detection of a neoplasm in a patient comprising an anti-endocan antibody.

26-75. (Cancelled)